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Patent
Attorney's Docket No. OPF 22.02

**UTILITY PATENT
APPLICATION TRANSMITTAL LETTER**

Box PATENT APPLICATION

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Enclosed for filing is the utility patent application of Emily L. BELLHOUSE, Stuart G. WEEKES, and Graeme L. J. PATERSON for PARTICLE DELIVERY DEVICE, claiming priority to US provisional application serial no. 60/117,806, filed January 29, 1999.

Also enclosed are:

5 sheet(s) of [] formal informal drawing(s); and

Other: Title Page and a return receipt postcard.

The declaration (unexecuted) of the inventor(s) also is enclosed.

The filing fee has been calculated as follows:

CLAIMS					
	NO. OF CLAIMS		EXTRA CLAIMS	RATE	FEE
Basic Application Fee					\$ 760.00
Total Claims	9	MINUS 20=	0	x \$18 =	.00
Independent Claims	2	MINUS 3 =	0	x \$78 =	.00
If multiple dependent claims are presented, add \$260.00					0
Total Application Fee					760.00
Add Assignment Recording Fee of \$40.00 if Assignment document is enclosed					0

Utility Patent Application Transmittal Letter
Attorney's Docket No.: OPF 22.02
Page 2

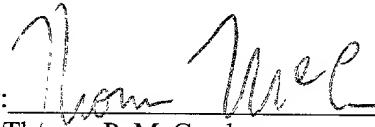
TOTAL APPLICATION FEE DUE	\$760.00
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No fees are enclosed.

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Respectfully submitted,

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OPF 22.02

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Application for U.S. Letters Patent, Entitled:

PARTICLE DELIVERY DEVICE

claiming priority to U.S. provisional application serial no. 60/117,806, filed January 29, 1999.

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PARTICLE DELIVERY DEVICE

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Cross-Reference to Related Application

This application is related to U.S. provisional patent application serial number 60/117,806, filed 29 January 1999, from which application priority is claimed pursuant to 35 U.S.C. §119(e)(1) and which application is incorporated herein by reference in its entirety.

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Technical Field

The invention relates generally to a needleless syringe for use in delivery of particles of a therapeutic agent to a target surface. More particularly, the invention pertains to a reusable needleless syringe system that uses a replaceable cartridge unit having a sealed chamber, a source of particles of a therapeutic agent, and a source of a driving gas contained at super-atmospheric pressure. The invention further pertains to a reusable needleless syringe system that uses a replaceable nozzle assembly.

Background

In commonly-owned U.S. Patent No. 5,630,796 to Bellhouse et al., a non-invasive delivery system is described that entails the use of a needleless syringe. The syringe is used for transdermal delivery of powdered therapeutic compounds and compositions to skin, muscle, blood or lymph. The syringe can also be used in conjunction with surgery to deliver therapeutics to organ surfaces, solid tumors and/or to surgical cavities (e.g., tumor beds or cavities after tumor resection).

The needleless syringe is constructed as an elongate tubular nozzle, typically having a rupturable membrane initially closing the passage through the nozzle adjacent to the upstream end of the nozzle. Particles of a powdered therapeutic agent are located adjacent to the membrane. The therapeutic agent is delivered using an energizing

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means which applies a gaseous pressure to the upstream side of the membrane that is sufficient to burst the membrane, thereby producing a supersonic gas flow through the nozzle. The supersonic gas flow produced in the nozzle entrains the particles and accelerates them to suitable velocities for transdermal delivery into tissue such as skin or muscle.

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Summary of the Invention

It is a primary object of the invention to provide a reusable needleless syringe device that employs one or more replaceable components. Accordingly, in one aspect of the invention, a replaceable cartridge is provided for use in a needleless syringe. The replaceable cartridge is formed from a housing having a top end and a bottom end, wherein the bottom end has an inlet aperture and an outlet aperture that are side-by-side and in fluid communication with each other by way of an internal gas passageway which extends through the housing. The cartridge further includes a particle container containing a suitable amount of particles for a single therapeutic or diagnostic particle administration. The particle container is arranged within the housing and positioned over the outlet. The cartridge also contains a source of compressed gas, again suitable for powering a single therapeutic or diagnostic particle administration. The source of compressed gas is arranged in sealable communication with the inlet, wherein the source includes an exterior portion which extends away from the housing and an interior portion which extends through the inlet and partially into the gas passageway such that a breach can be made to the interior portion in order to release compressed gas from the source and into the gas passageway.

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In one configuration of the replaceable cartridge, the internal gas passageway is substantially in the shape of a U. In addition, the replaceable cartridge can accommodate an inserted expansion chamber at the downstream end of the gas passageway, or the housing can be molded or otherwise constructed to include an expansion chamber portion within the gas passageway. A filter element can likewise be included within the gas passageway.

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In another aspect of the invention, a replaceable nozzle assembly is provided for use in a needleless syringe. The replaceable nozzle assembly includes an elongate particle acceleration nozzle having an upstream end, a downstream end and a gas passageway extending between the upstream and downstream ends. The gas passageway has an upper convergent section, a lower divergent section, and a throat section connecting the convergent and divergent sections. An outwardly projecting, disk-shaped annular flange is provided at the upstream terminus of the nozzle, which flange provides an external shoulder. The nozzle assembly further includes a cylindrical filter element which fits over and substantially surrounds the elongate particle acceleration nozzle. The filter element is separately replaceable from the nozzle. The filter element rests upon the external shoulder provided by the flange. The filter element includes a filter medium which provides a barrier against fine particles which, having insufficient momentum, may rebound from the target surface during or shortly after a particle delivery operation has been carried out.

In a still further aspect of the invention, a reusable needleless syringe device is provided. The reusable needleless syringe contains either or both the replaceable cartridge and the replaceable nozzle assembly of the present invention.

It is an advantage of the invention that a reusable needleless syringe can be provided which allows for multiple particle delivery operations to be carried out. It is a further advantage of the invention that the replaceable cartridges and nozzle assemblies of the present invention can be easily mass produced, packaged, stored, handled and then disposed of, significantly reducing the per-shot cost of operating a needleless syringe device.

These and other objects, aspects, embodiments and advantages of the present invention will readily occur to those of ordinary skill in the art in view of the disclosure herein.

Brief Description of the Figures

Figure 1 is a cross-sectional view of a reusable needleless syringe device

containing a replaceable nozzle assembly and a replaceable cartridge, both of which replaceable components are constructed according to the present invention.

Figure 2 is a pictorial representation of the reusable needleless syringe device of Figure 1 shown in partial cut-away to illustrate the working components of the device.

5 Figure 3 is a cross-sectional view of a replaceable cartridge constructed according to the invention.

Figure 4 is a side plan view of the replaceable cartridge of Figure 3.

Figure 5 is a top plan view of the replaceable cartridge of Figure 3.

Figure 6 is an exploded view of the replaceable cartridge of Figure 3.

10 Figure 7 is a pictorial representation depicting the replaceable cartridge of Figure 3 being inserted into a reusable needleless syringe device similar to the device of Figures 1 and 2.

15 Figure 8 is a side plan view of a replaceable nozzle assembly constructed according to the present invention, wherein the assembly is shown in partial exploded view.

Figure 9 is a pictorial representation depicting the replaceable nozzle assembly of Figure 8 being inserted into a reusable needleless syringe device similar to the device of Figures 1 and 2.

20 Detailed Description of the Preferred Embodiments

Before describing the present invention in detail, it is to be understood that this invention is not limited to particularly exemplified device structures as such may, of course, vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments of the invention only, and is not intended to be limiting.

25 All publications, patents and patent applications cited herein, whether *supra* or *infra*, are hereby incorporated by reference in their entirety.

It must be noted that, as used in this specification and the appended claims, the singular forms "a," "an" and "the" include plural referents unless the content clearly

dictates otherwise. Thus, for example, reference to "a therapeutic agent" includes a mixture of two or more such agents, reference to "a gas" includes mixtures of two or more gases, and the like.

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A. Definitions

Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which the invention pertains. Although a number of methods and materials similar or equivalent to those described herein can be used in the practice of the present invention, the preferred materials and methods are described herein.

10 In describing the present invention, the following terms will be employed, and are intended to be defined as indicated below.

15 The terms "needleless syringe," and "needleless syringe device," as used herein, expressly refer to a particle delivery system that can be used to deliver particles into and/or across tissue, wherein the particles have an average size ranging from about 0.1 to 250 μm , preferably about 10-70 μm . Particles larger than about 250 μm can also be delivered from these devices, with the upper limitation being the point at which the size of the particles would cause untoward pain and/or damage to the target tissue. The particles are delivered at high velocity, for example at velocities of at least about 150 m/s or more, and more typically at velocities of about 250-300 m/s or greater. Such needleless syringe devices were first described in commonly-owned U.S. Patent No. 20 5,630,796 to Bellhouse et al., incorporated herein by reference, and have since been described in commonly owned International Publication Nos. WO 96/04947, WO 12513, and WO 96/20022, all of which publications are also incorporated herein by reference. These devices can be used in the transdermal delivery of a therapeutic agent through skin or mucosal tissue, either *in vitro* or *in vivo* (*in situ*); or the devices can be used in the transdermal delivery of generally inert particles for the purpose of non- or minimally invasive sampling of an analyte from a biological system. Since the term 25 only relates to devices which are suitable for delivery of particulate materials, devices

such as liquid-jet injectors are expressly excluded from the definition of a "needleless syringe."

The term "transdermal" delivery captures intradermal, transdermal (or "percutaneous") and transmucosal administration, i.e., delivery by passage of a therapeutic agent into and/or through skin or mucosal tissue. See, e.g., *Transdermal Drug Delivery: Developmental Issues and Research Initiatives*, Hadgraft and Guy (eds.), Marcel Dekker, Inc., (1989); *Controlled Drug Delivery: Fundamentals and Applications*, Robinson and Lee (eds.), Marcel Dekker Inc., (1987); and *Transdermal Delivery of Drugs*, Vols. 1-3, Kydonieus and Berner (eds.), CRC Press, (1987). Aspects of the invention which are described herein in the context of "transdermal" delivery, unless otherwise specified, are meant to apply to intradermal, transdermal and transmucosal delivery. That is, the devices, systems, and methods of the invention, unless explicitly stated otherwise, should be presumed to be equally applicable to intradermal, transdermal and transmucosal modes of delivery.

As used herein, the terms "therapeutic agent" and/or "particles of a therapeutic agent" intend any compound or composition of matter which, when administered to an organism (human or animal) induces a desired pharmacologic, immunogenic, and/or physiologic effect by local and/or systemic action. The term therefore encompasses those compounds or chemicals traditionally regarded as drugs, vaccines, and biopharmaceuticals including molecules such as proteins, peptides, hormones, biological response modifiers, nucleic acids, gene constructs and the like. More particularly, the term "therapeutic agent" includes compounds or compositions for use in all of the major therapeutic areas including, but not limited to, adjuvants, anti-infectives such as antibiotics and antiviral agents; analgesics and analgesic combinations; local and general anesthetics; anorexics; antiarthritics; antiasthmatic agents; anticonvulsants; antidepressants; antigens, antihistamines; anti-inflammatory agents; antinauseants; antineoplastics; antipruritics; antipsychotics; antipyretics; antispasmodics; cardiovascular preparations (including calcium channel blockers, beta-blockers, beta-agonists and antiarrhythmics); antihypertensives; diuretics; vasodilators;

central nervous system stimulants; cough and cold preparations; decongestants; diagnostics; hormones; bone growth stimulants and bone resorption inhibitors; immunosuppressives; muscle relaxants; psychostimulants; sedatives; tranquilizers; proteins peptides and fragments thereof (whether naturally occurring, chemically synthesized or recombinantly produced); and nucleic acid molecules (polymeric forms of two or more nucleotides, either ribonucleotides (RNA) or deoxyribonucleotides (DNA) including both double- and single-stranded molecules, gene constructs, expression vectors, antisense molecules and the like).

Particles of a therapeutic agent, alone or in combination with other drugs or agents, are typically prepared as pharmaceutical compositions which can contain one or more added materials such as carriers, vehicles, and/or excipients. "Carriers," "vehicles" and "excipients" generally refer to substantially inert materials which are nontoxic and do not interact with other components of the composition in a deleterious manner. These materials can be used to increase the amount of solids in particulate pharmaceutical compositions. Examples of suitable carriers include water, silicone, gelatin, waxes, and like materials. Examples of normally employed "excipients," include pharmaceutical grades of dextrose, sucrose, lactose, trehalose, mannitol, sorbitol, inositol, dextran, starch, cellulose, sodium or calcium phosphates, calcium sulfate, citric acid, tartaric acid, glycine, high molecular weight polyethylene glycols (PEG), and combinations thereof. In addition, it may be desirable to include a charged lipid and/or detergent in the pharmaceutical compositions. Such materials can be used as stabilizers, anti-oxidants, or used to reduce the possibility of local irritation at the site of administration. Suitable charged lipids include, without limitation, phosphatidylcholines (lecithin), and the like. Detergents will typically be a nonionic, anionic, cationic or amphoteric surfactant. Examples of suitable surfactants include, for example, Tergitol® and Triton® surfactants (Union Carbide Chemicals and Plastics, Danbury, CT), polyoxyethylenesorbitans, e.g., TWEEN® surfactants (Atlas Chemical Industries, Wilmington, DE), polyoxyethylene ethers, e.g., Brij, pharmaceutically acceptable fatty acid esters, e.g., lauryl sulfate and salts thereof (SDS), and like

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materials.

The term "analyte" is used herein in its broadest sense to denote any specific substance or component that one desires to detect and/or measure in a physical, chemical, biochemical, electrochemical, photochemical, spectrophotometric, polarimetric, colorimetric, or radiometric analysis. A detectable signal can be obtained, either directly or indirectly, from such a material. In some applications, the analyte is a physiological analyte of interest (e.g., a physiologically active material), for example glucose, or a chemical that has a physiological action, for example a drug or pharmacological agent.

As used herein, the term "sampling" means extraction of a substance from any biological system across a membrane, generally across skin or tissue. The membrane can be natural or artificial, and is generally animal in nature, such as natural or artificial skin, blood vessel tissue, intestinal tissue, and the like. A "biological system" thus includes both living and artificially maintained systems.

B. General Methods

In one embodiment, a replaceable cartridge for use with a reusable needleless syringe is provided. The replaceable cartridge contains both a source of particles and a source of compressed gas, and is intended as a single-use disposable cartridge. In another embodiment, a replaceable nozzle assembly for use with a reusable needleless syringe is provided. The replaceable nozzle assembly includes a supersonic particle acceleration nozzle and a filter element which surrounds the acceleration nozzle. In a still further embodiment, a reusable needleless syringe is provided with contains either one or both of the above-described replaceable components.

Needleless syringe devices for delivering particles were first described in commonly owned U.S. Patent No. 5,630,796 to Bellhouse et al., incorporated herein by reference. Although a number of specific device configurations are now available, such devices are typically provided as a pen-shaped instrument containing, in linear order moving from top to bottom, a gas cylinder, a particle cassette or package, and a

supersonic nozzle with an associated silencer medium. An appropriate powder is sealed within a particle cassette, e.g., a container formed by two rupturable polymer membranes that are heat-sealed to a washer-shaped spacer to form a self-contained sealed unit. The membrane material is selected to achieve a specific mode of opening and burst pressure that dictates the conditions at which the supersonic flow is initiated. In operation, the device is actuated to release the compressed gas from the cylinder into an expansion chamber within the device. The released gas contacts the particle cassette and, when sufficient pressure is built up, suddenly breaches the cassette membranes sweeping the particles into the supersonic nozzle for subsequent delivery. The nozzle is designed to achieve a specific gas velocity and flow pattern to deliver a quantity of particles to a target surface of predefined area. The silencer is used to attenuate the noise of the "sonic boom" produced by the membrane rupture. Such devices are most commonly manufactured as a disposable unit, and the spent devices are intended to be discarded after a single particle delivery operation has been carried out. Accordingly, these devices are typically inexpensively manufactured from medical grade plastics using injection-molding techniques (with the exception of the gas cylinder which is typically deep-drawn from aluminum or some other suitable metal or metal alloy) to facilitate such single use economics.

The reusable needleless syringes of the present invention contain the same basic elements and function in the same basic manner as the above-described devices. However, these reusable devices are specially configured to accommodate multiple use by either medical personnel or consumers, and contain replaceable components such as a cartridge containing a fresh particle cassette with a gas cylinder, or a new nozzle unit. One particular reusable needleless syringe device is depicted in Figures 1 and 2. As can be seen, this particular reusable device is not arranged in the typical linear "pen-shaped" configuration, rather the device has been bent around into a U-shape to place the nozzle and gas cylinder into a "side-by-side" orientation, substantially shortening the overall length of the device.

Referring particularly to Figure 1, the reusable device 2 includes a replaceable

nozzle assembly generally indicated at 4 which has a supersonic particle acceleration nozzle. The particle acceleration nozzle has an upper convergent section 6, a lower divergent section 8, and a throat section 10 connecting the convergent and divergent sections. The particle acceleration nozzle terminates at its downstream end within a spacer shroud 12 that is provided by the reusable device 2. The shroud is intended to be pressed against a target surface to create a suitable seal and to space the nozzle at a preselected distance from the surface to optimize particle delivery conditions. The outside of the nozzle assembly 4 is surrounded by a filter element 14 that is intended to capture and contain any particle "fines" that may rebound from the target surface and be dispersed within the spacer shroud and nozzle assembly immediately after delivery. The shroud is further intended to capture the shockwave that is reflected from the target surface, which shockwave can then travel radially outwards from the shroud through appropriate vents and pass into a silencer chamber 16 for sound attenuation. As will be described in greater detail herein below, the filter element 14 can be easily disconnected from the rest of the nozzle assembly 4 and thus replaced separately.

The reusable device 2 further includes a replaceable cartridge 18 which comprises a housing 20, a source of compressed gas 22, and a particle container 24 which contains particles. The housing 20 provides an enclosed structure having an internal gas passageway 26 which connects an inlet aperture 28 and an outlet aperture 30 which are disposed on the same bottom side of the housing. The source of compressed gas 22 is inserted into the inlet aperture 28 such that a top portion of the source extends partially into the internal gas passageway 26. Insertion of the gas source, in this case, a gas cylinder, into the inlet aperture closes off the inlet and provides a gas-tight seal. Similarly, the particle container 24 is positioned over and blocks off the outlet aperture 30 also creating a gas-tight seal. In this way, the replaceable cartridge 18 is an entirely self-contained, integral component which provides both the particles and the source of motive force necessary for a single operation of the reusable needleless syringe device. The particle container 24 can comprise any suitable structure which can effectively contain a metered amount of

5 particles. However, in the particular embodiment depicted in Figure 1, the particle container is a capsule formed from two comparatively weak, rupturable membranes which are attached to a spacer ring to establish a self-contained unit as described in U.S. Patent No. 5,630,796. The rupturable membranes are formed from a suitable polymer material, for example polycarbonates (e.g., MacrafolTM), MYLAR[®] or other like materials. Both the membrane material and the thickness of the membrane material are selected to provide for a specific mode of opening and burst pressure as also previously described.

10 Referring now to Figures 1 and 2, the reusable device is operated by creating a breach in the top portion of the gas source 22 such that the compressed gas can escape from the source and enter into the internal gas passageway 26. In the particular embodiment depicted in Figures 1 and 2, the breach is created by the action of a actuation ram or pin 32 which is used to snap off a frangible tip 34 from the inner portion of the gas source. In this manner, the compressed gas flows into the internal gas passageway 26 where it is initially retained by the upstream membrane of the particle container 24. After sufficient pressure has built up within the gas passageway, the rupturable membranes of the particle container are burst, allowing the gas to sweep through the container and entrain the particles within the gas flow entering the supersonic particle acceleration nozzle of the nozzle assembly 4. A number of 15 alternative actuation mechanisms can, of course be used to create the breach in the gas source. For example, a sharp pin or needle can be used to pierce a hole in the gas source or rupture a membrane or other relatively weak portion of the gas source. Alternatively, a trigger mechanism can be used to open a valve which closes off the gas source. These and other suitable actuation schemes and mechanisms will readily occur to the 20 ordinarily skilled artisan upon reading the instant specification. After the device has been operated, the spent cartridge 18 can be removed from the reusable device 2 and replaced with a fresh cartridge which contains a new gas source and particle container. In like manner, the nozzle assembly 4 can be removed, and either the filter element 14, 25 or the entire nozzle assembly can be replaced with fresh components.

Figures 3-7 depict the construction and assembly of a replaceable cartridge 18 which is suitable for use in the reusable needleless syringe devices of the present invention. Referring particularly to Figures 3 and 6, the cartridge housing 20 is roughly in the shape of a D and has inlet and outlet apertures, 28 and 30, respectively, positioned on the bottom end thereof. The housing can be made from any suitably resilient material, preferably from a medical-grade plastic which may be injection-molded into any suitable configuration.

During assembly, a fresh gas source 22 is inserted into the inlet aperture 28. The gas source can have a flange 36 which rests against a corresponding shoulder within the housing in order to position the tip 34 within the gas passageway 26 as previously described. When the cartridge is fully assembled, this flange helps provide for a gas-tight seal. The gas source 22 can be any suitable container of compressed air or gas, and typically will be charged at about 10-100 bar or more, more typically at about 25-60 bar. The top end of the cartridge housing 20 is hollow in order to establish an intermediate portion of the gas passageway, and is typically somewhat pinched and narrow as best seen in the top view provided by Figure 5. The gas passageway 26 is substantially in the shape of a U. An optional internal filter element 38 can be arranged within the internal gas passageway 26 in order to prevent fragments or bits from the breached gas source from entering into the nozzle assembly. The internal filter element is preferably arranged at or near the outlet aperture 30, and can be made from any suitable filtering material, for example a stainless steel mesh or fabric material that allows free passage of the compressed gas but retains any fragments or particles which could enter into the gas passageway during operation of the device. The internal filter element 38 can rest upon an internal shoulder provided within the housing, or can be positioned and retained by any suitable means.

The cartridge housing 20 can be manufactured to provide an expansion chamber portion within the gas passageway, at a location adjacent to the outlet aperture 30. In this regard, the expansion chamber portion is configured to have an increased cross-sectional area relative to the intermediate portion of the passageway, and thus allows for

the creation of an increased pressure front immediately above the particle source 24 in order to facilitate operation of the device. However, it is preferred that an expansion chamber insert 40 be used to provide this function, wherein the insert can be a simple molded annular part having a suitable convergent inner geometry.

Accordingly, during assembly, the internal filter element can be dropped into the outlet aperture 30 and rests upon an internal shoulder. The expansion chamber insert 40 can then be inserted, and the particle container 24 finally added to block off the outlet aperture. A bottom casement 42 can then be snapped into place or otherwise attached to the housing 20 in order to hold the various components together as a fully assembled, gas-tight replaceable cartridge. As can be seen in Figure 3, the bottom casement 42 can optionally have an outlet aperture portion 48 which provides an annular female lip at the terminus of the outlet aperture 30 in the assembled cartridge. This annular lip accepts and closely engages a corresponding annular male flange provided at the top of a nozzle assembly, thus positioning the particle container immediately over the entrance of the particle acceleration nozzle and providing a tight seal between the two elements.

Referring particularly to Figures 3 and 4, when the cartridge has been assembled, an interior portion of the gas source 22 extends partially into the gas passageway 26, and an exterior portion of the gas source extends away from the housing 20. Part of the interior portion of the gas source is comprised of a frangible, elongate tip 34 which is positioned in the assembled cartridge so that it is adjacent to an actuation aperture 44. The actuation aperture is sized to allow an actuation ram or pin to travel into the housing and break off the frangible tip during actuation of the device. The actuation aperture 44 is sealed off by a pierceable foil or seal 44 which is applied to the outside of the housing as seen in Figure 3. This seal provides for an intuitive indicator of when a cartridge has been spent, since the seal on a used cartridge will be breached.

The fully assembled replaceable cartridge is inserted into a reusable needleless syringe device as follows. Referring particularly to Figure 7, the back cover 54 of a reusable needleless syringe device 52 is opened to reveal a cavity 56 and the top end of a replaceable nozzle assembly 4. The exterior portion of the gas source 22 fits into the

cavity 56 and the outlet aperture 30 couples with the top of the nozzle assembly 4. The cover 54 is then closed to provide a fully ready and operational device.

Other embodiments of the replaceable cartridge of the present invention are those which can be used with the more standard, linear "pen-shaped" devices such as the needleless syringe devices disclosed in U.S. Patent No. 5,630,796 to Bellhouse et al. Accordingly, in such alternative embodiments, the cartridge will have an overall cylindrical configuration, with the various components being provided in a linear arrangement. Such cartridges can be thought of as a shotgun shell, wherein a source of motive force (the gas source) is positioned behind a particle container and both elements are encased by an easy to handle outer housing. One particular arrangement comprises a housing having, in linear arrangement in the order listed, an upstream gas source (e.g., a cartridge containing compressed gas or air), an intermediate gas passageway including an expansion chamber, and a downstream particle container. Optional elements, such as an internal filter or an inserted expansion chamber, can be included as described above for the side-by-side cartridge. The gas source can be fully contained within the housing or, alternatively, have an interior portion extending partially into the gas passageway and an external portion extending away from the housing as also described above. In this manner, a breach can be made to the interior portion of the gas source in order to actuate a particle delivery operation.

Figures 8 and 9 depict the construction and assembly of a replaceable nozzle assembly 4 which is suitable for use in the reusable needleless syringe devices of the present invention. Referring particularly to Figure 8, the replaceable nozzle assembly 4 comprises an elongate particle acceleration nozzle 60 and a cylindrical filter element 14 which is inserted over and substantially surrounds the particle acceleration nozzle. The particle acceleration nozzle 60 can have one or more outwardly projecting fins 62 depending from the exterior surface thereof, which fins may provide structural rigidity to the nozzle itself and/or establish a support for the cylindrical filter element. The passageway through the nozzle has an upstream convergent section, a downstream divergent section, and a throat section interposed between the convergent and divergent

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sections. The particle acceleration nozzle can be formed from any suitably resilient material which can be machined and/or molded to provide both the desired external features, e.g., the outwardly projecting fins 62 and the desired external features, e.g., the convergent, divergent nozzle geometries. It is preferred that the nozzle is injection-molded from a suitable medical grade of plastic. The particle acceleration nozzle further comprises an outwardly projecting disc-shaped flange 64 depending from its upstream end to provide an external shoulder. The flange 64 may optionally have a finger or clip 66 extending therefrom in a downstream direction and providing a point of engagement for the filter element 14 as described herein below. The flange 64 may also have a grip or tab 68 extending therefrom in an upstream direction, which grip or tab provides an intuitive element for grasping the replaceable nozzle assembly 4 and removing the same when the assembly is properly positioned in a reusable needleless syringe.

The filter element 14 comprises a cylindrical carrier moulding 70 which is sized to fit around and substantially surround the elongate particle acceleration nozzle 60 along most of its length. The carrier moulding 70 is preferably constructed as an open structure, for example any honeycomb, scaffold, lattice or otherwise open-frame construction which provides for as much open area about its periphery as possible. The carrier moulding can be formed from any suitable material such as a metal, plastic, rigid paper, cardboard or pressed-fiber material. A filter medium 72 is wrapped around the carrier molding and establishes a barrier against reflected particles and/or particle "fines" that, having insufficient momentum to penetrate the intended target, may rebound from the target surface during or just after particle delivery from a needleless syringe. This helps prevent such particles or particle fines from entering into inappropriate parts of the reusable device, particularly from entering into the silencing medium generally used with needleless syringe devices which silencing medium comprises an open-cell foam material. The filter medium is selected to have a very small pore size and can thus be formed from a paper, fiber, or paper/fiber filter sheet similar to a coffee filter. If desired, the carrier moulding 70 can have a notch or other

fastener mechanism 74 that cooperates with the corresponding finger or clip 66 on the nozzle flange 64. The replaceable nozzle assembly is put together by sliding the cylindrical carrier molding over the outside of the nozzle 60, generally using the fins 62 to guide and support the carrier until it is brought to rest against the shoulder provided by the flange 64. The two elements can be held together as a complete nozzle assembly by way of establishing a tight fit tolerance between the outside dimension of the particle acceleration nozzle 60, e.g., the outermost edges of the fins 62, and the inside dimension of the carrier moulding 70. Alternatively or additionally, the two elements can be held together by a latch or fastening mechanism provided by one or both of the carrier moulding and the nozzle, e.g., the corresponding structures 74 and 66 on each respective element. Removal of the filter element 14 from the particle acceleration nozzle 60 is carried out by pulling the two elements apart in the directions depicted by the right and left opposing arrows in Figure 8. This allows the filter element to be replaced separately from, and possibly at more frequent intervals than the particle acceleration nozzle in a reusable needleless syringe device.

The fully assembled replaceable nozzle assembly is inserted into a reusable needleless syringe device as follows. Referring now to Figure 9, the back cover 84 of a reusable needleless syringe device 82 is opened to reveal a cavity 86 which can accept and closely retain the nozzle assembly 4. If desired, a gasket 88 can be used to provide a tight seal between the flange 64 of the nozzle assembly and the case of the reusable needleless syringe device 82. The cover 84 is then closed to provide a fully ready and operational device. As will be understood by those skilled in the art upon reading the instant specification, the replaceable nozzle assembly of the present invention can be used with any reusable needleless syringe configuration, including the "side-by-side" configuration described herein, as well as the more traditional "pen-shaped" device as disclosed in U.S. Patent No. 5,630,796 to Bellhouse et al.

Accordingly, novel replaceable components for use with a reusable needleless syringe system have been described, as well as reusable needleless syringe devices which comprise such replaceable components. Although preferred embodiments of the

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subject invention have been described in some detail, it is understood that obvious variations can be made without departing from the spirit and the scope of the invention as defined by the appended claims.

We claim:

1. A replaceable cartridge for use in a needleless syringe, said cartridge comprising:

5 (a) a housing having a top end and a bottom end, wherein the bottom end has an inlet aperture and an outlet aperture in fluid communication with each other by way of an internal gas passageway which extends through said housing;

10 (b) a particle container containing particles, said container arranged within the housing and positioned over the outlet; and

15 (c) a source of compressed gas in sealable communication with the inlet, wherein said source of compressed gas includes an exterior portion which extends away from the housing and an interior portion which extends through the inlet and partially into the gas passageway such that a breach can be made to the interior portion in order to release compressed gas from the source and into the gas passageway.

20 2. The cartridge of claim 1, wherein said gas passageway is substantially in the shape of a U.

25 3. The cartridge of claim 1, wherein said gas passageway has an expansion chamber portion of increased cross-sectional area, said expansion chamber having an upstream end and a downstream end and positioned within the housing just above the outlet such that the particle container can be arranged immediately between the downstream end of the expansion chamber and the outlet.

4. The cartridge of claim 1 further comprising a filter element positioned within the gas passageway between the inlet and the outlet apertures.

5. The cartridge of claim 4, wherein the filter element is positioned at the upstream end of the expansion chamber.

6. A replaceable nozzle assembly for use in a needleless syringe, said nozzle assembly comprising:

(a) an elongate particle acceleration nozzle having an upstream end, a downstream end and a passageway extending between the upstream and downstream ends, wherein said passageway has an upper convergent section, a lower divergent section, and a throat section connecting the convergent and divergent sections, and further wherein said nozzle has an outwardly projecting annular flange depending from its upstream end to provide an external shoulder; and

(b) a cylindrical filter medium which fits over and substantially surrounds the elongate particle acceleration nozzle and rests upon the external shoulder provided by the flange.

7. A needleless syringe device comprising the replaceable cartridge of claim 1.

8. The needleless syringe device of claim 7 further comprising the replaceable nozzle assembly of claim 6.

9. A needleless syringe device comprising the replaceable nozzle assembly of claim 6.

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PARTICLE DELIVERY DEVICE

Abstract of the Disclosure

Replaceable components for use with a needleless syringe delivery device are provided. One replaceable component is a cartridge having a source of particles and a source of compressed gas. Another replaceable component is a nozzle assembly having a particle acceleration nozzle and a coaxial filter element. Reusable needleless syringe devices which include these replaceable components are also provided.

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FIGURE 1

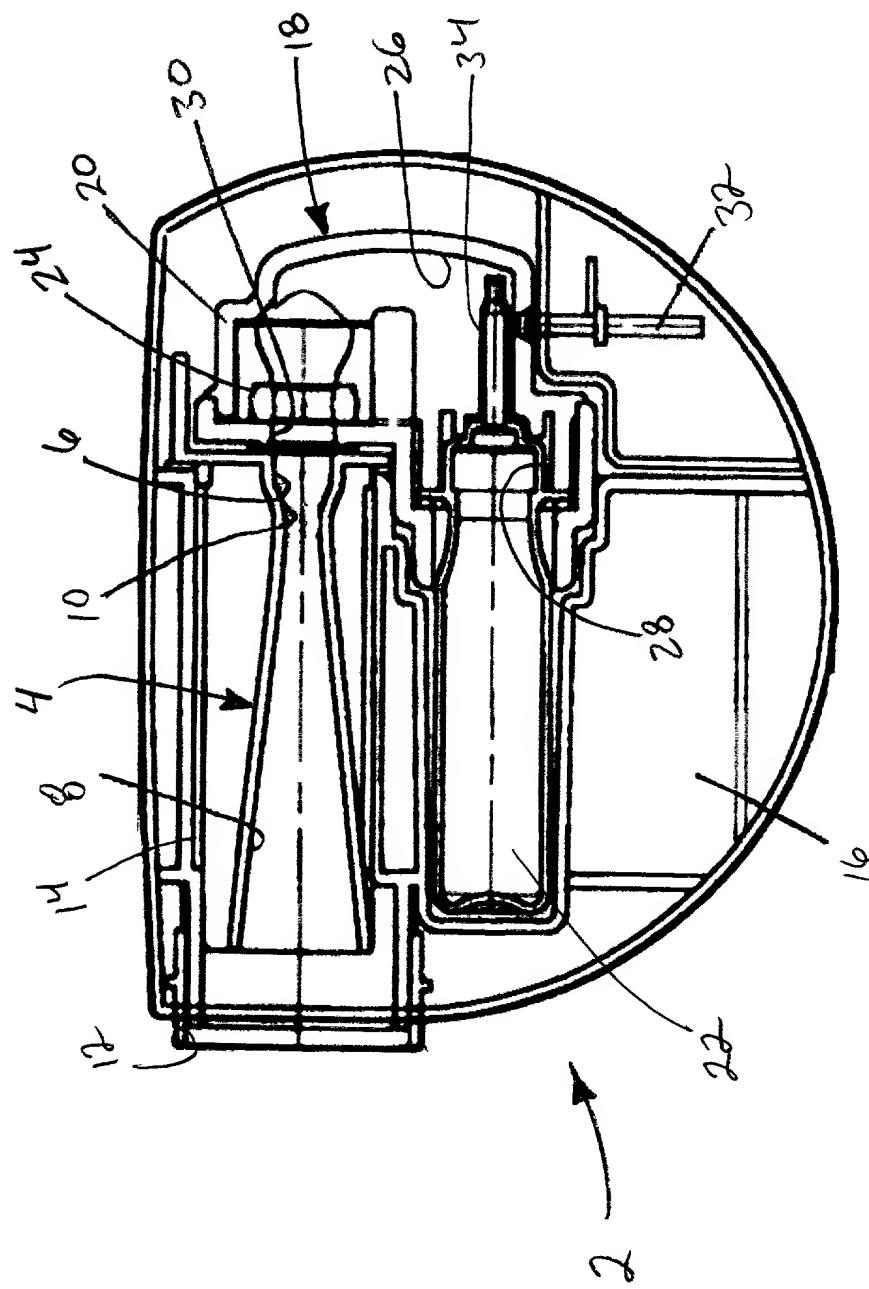
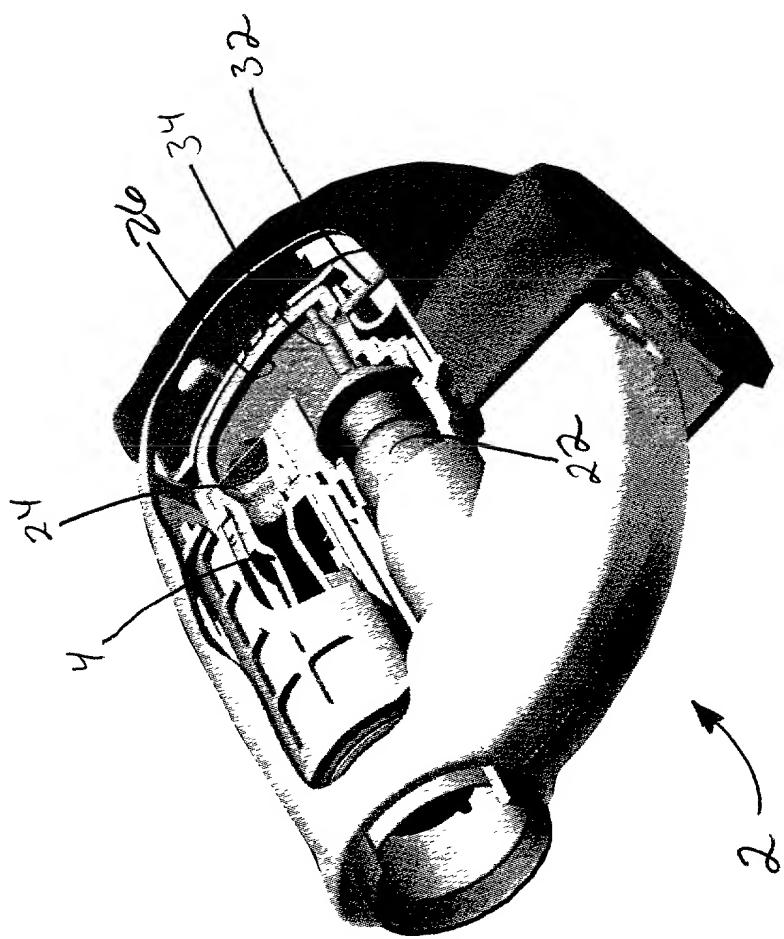


FIGURE 2



Patent Drawing of a Safety Belt Buckle

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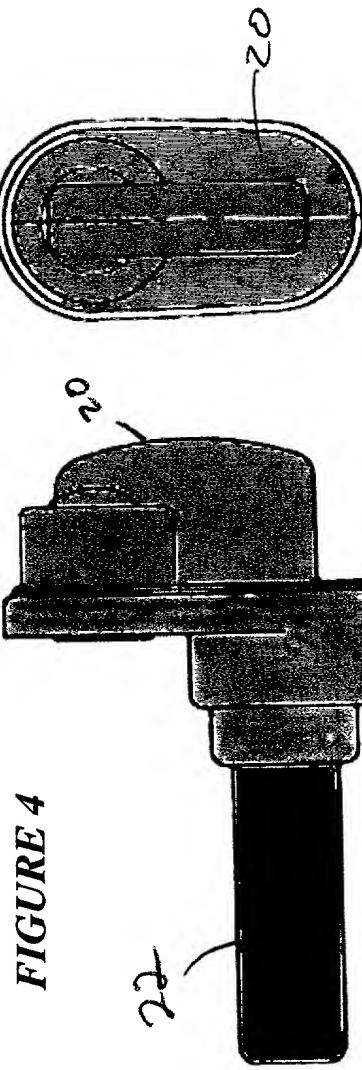


FIGURE 4

FIGURE 5

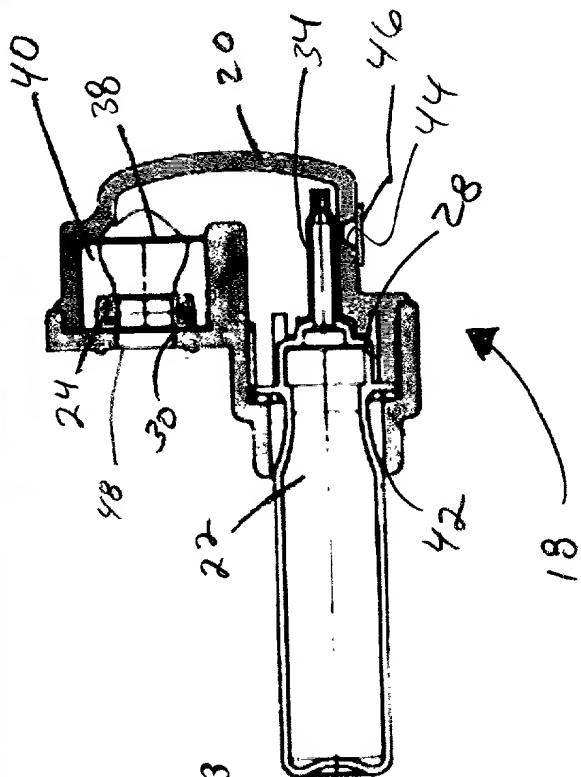


FIGURE 3

FIGURE 6

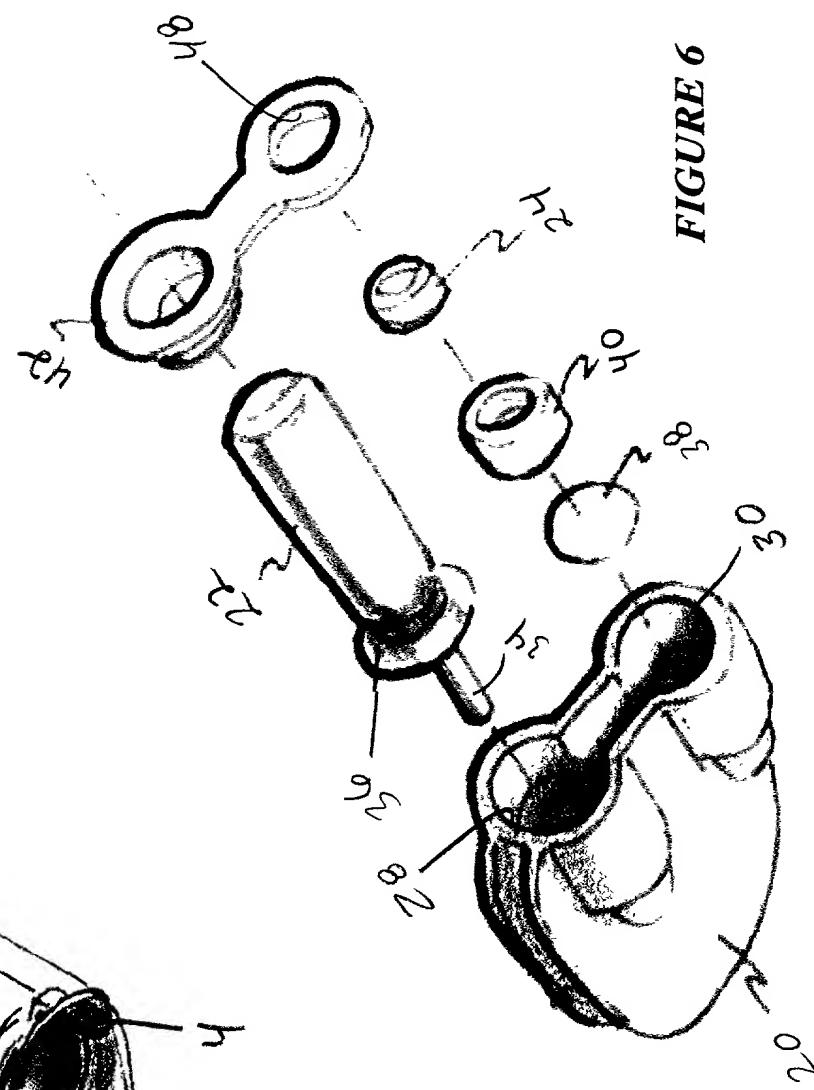
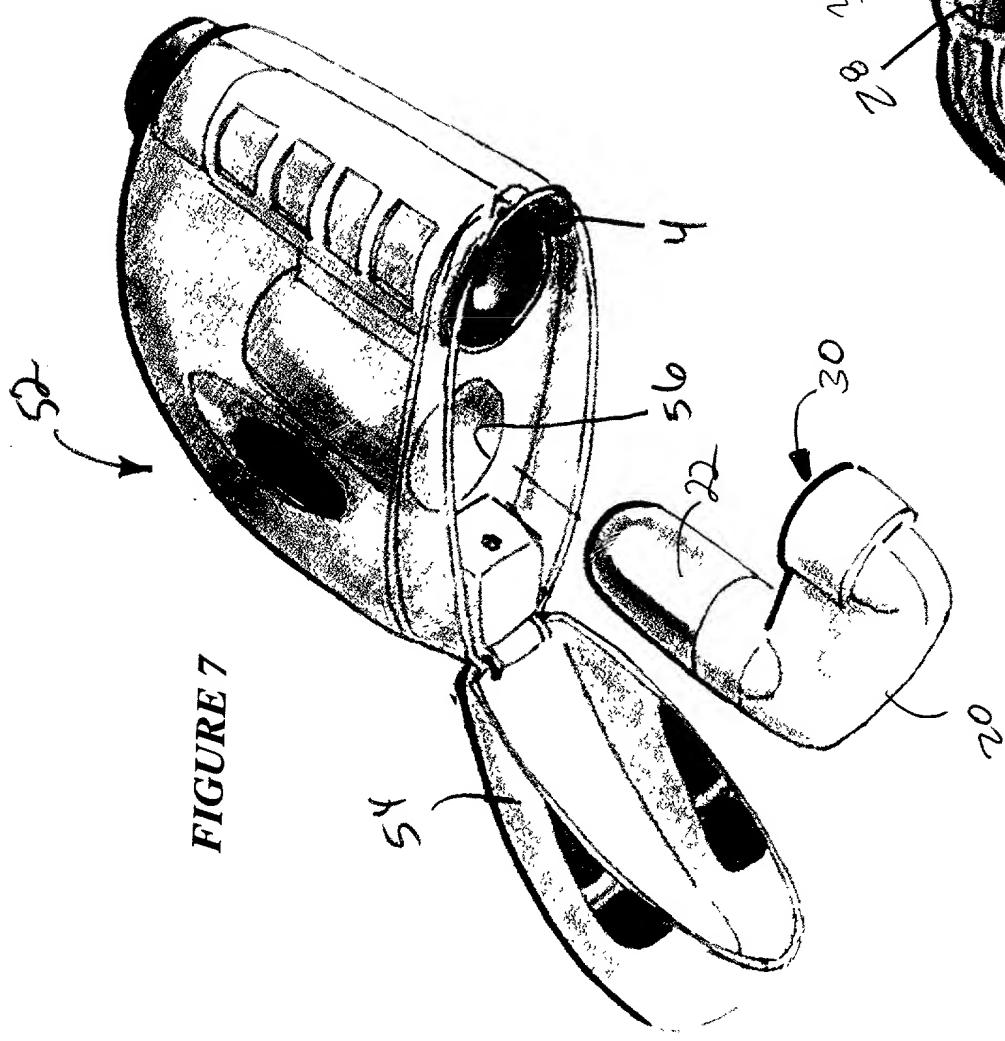
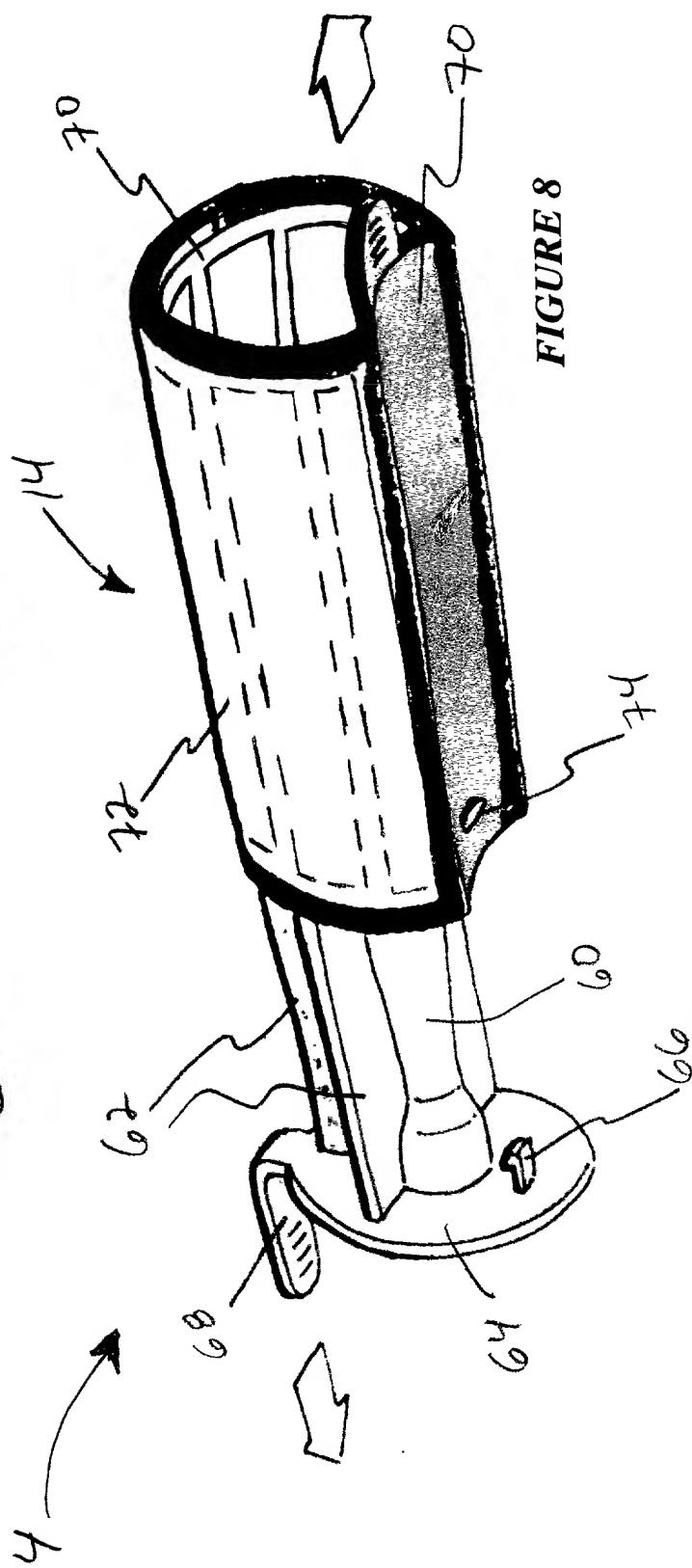
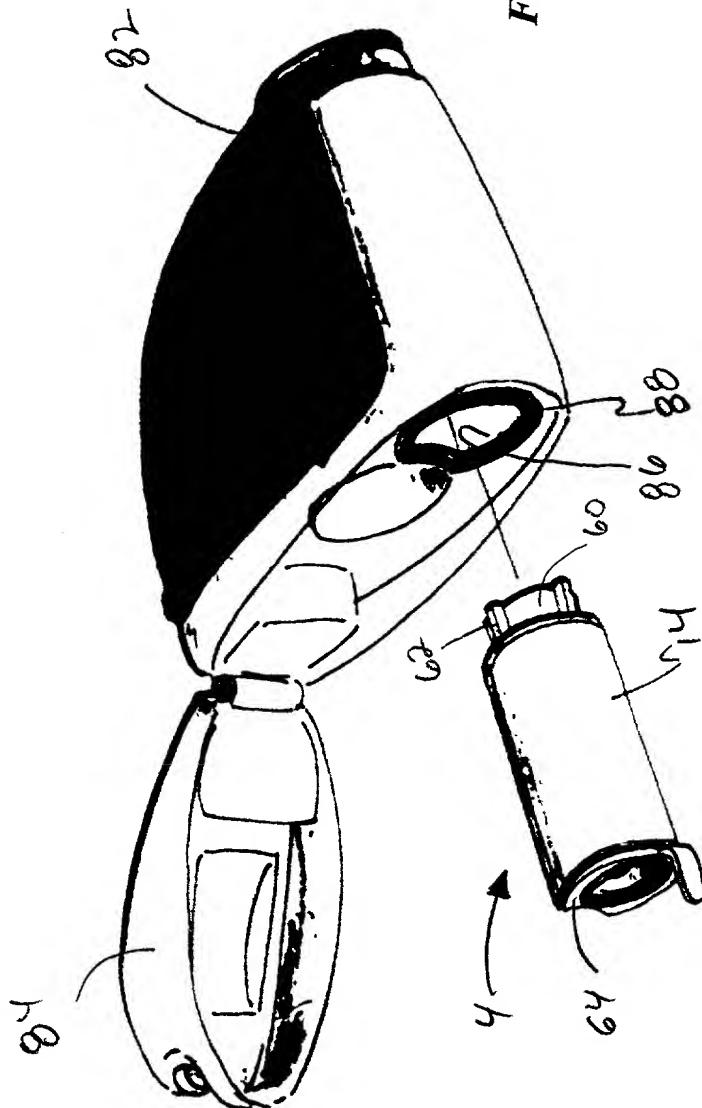


FIGURE 7





COMBINED DECLARATION AND POWER OF ATTORNEY
FOR UTILITY PATENT APPLICATION

AS A BELOW-NAMED INVENTOR, I HEREBY DECLARE THAT:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if more than one name is listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled: PARTICLE DELIVERY DEVICE the specification of which

X is attached hereto
____ was filed on _____, and assigned Serial No. _____.

I HAVE REVIEWED AND UNDERSTAND THE CONTENTS OF THE ABOVE-IDENTIFIED SPECIFICATION, INCLUDING THE CLAIMS, AS AMENDED BY ANY AMENDMENT REFERRED TO ABOVE.

I acknowledge and understand that I am an individual who has a duty to disclose information which is material to the patentability of the claims of this application in accordance with Title 37, Code of Federal Regulations, §§ 1.56(a) and (b) which state:

(a) A patent by its very nature is affected with a public interest. The public interest is best served, and the most effective patent examination occurs when, at the time an application is being examined, the Office is aware of and evaluates the teachings of all information material to patentability. Each individual associated with the filing and prosecution of a patent application has a duty of candor and good faith in dealing with the Office, which includes a duty to disclose to the Office all information known to that individual to be material to patentability as defined in this section. The duty to disclose information exists with respect to each pending claim until the claim is canceled or withdrawn from consideration, or the application becomes abandoned. Information material to the patentability of a claim that is canceled or withdrawn from consideration need not be submitted if the information is not material to the patentability of any claim remaining under consideration in the application. There is no duty to submit information which is not material to the patentability of any existing claim. The duty to disclose all information known to be material to patentability is deemed to be satisfied if all information known to be material to patentability of any claim issued in a patent was cited by the Office or submitted to the Office in the manner prescribed by §§ 1.97(b)-(d) and 1.98. However, no patent will be granted on an application in connection with which fraud on the Office was practiced or attempted or the duty of disclosure was violated through bad faith or intentional misconduct. The Office encourages applicants to carefully examine:

(1) prior art cited in search reports of a foreign patent office in a counterpart application, and

(2) the closest information over which individuals associated with the filing or prosecution of a patent application believe any pending claim patentably defines, to make sure that any material information contained therein is disclosed to the Office.

(b) Under this section, information is material to patentability when it is not cumulative to information already of record or being made of record in the application, and

(1) It establishes, by itself or in combination with other information, a *prima facie* case of unpatentability of a claim; or

(2) It refutes, or is inconsistent with, a position the applicant takes in:

- (i) Opposing an argument of unpatentability relied on by the Office, or
- (ii) Asserting an argument of patentability.

A *prima facie* case of unpatentability is established when the information compels a conclusion that a claim is unpatentable under the preponderance of evidence, burden-of-proof standard, giving each term in the claim its broadest reasonable construction consistent with the specification, and before any consideration is given to evidence which may be submitted in an attempt to establish a contrary conclusion of patentability.

I do not know and do not believe this invention was ever known or used in the United States of America before my or our invention thereof, or patented or described in any printed publication in any country before my or our invention thereof or more than one year prior to said application. This invention was not in public use or on sale in the United States of America more than one year prior to this application. This invention has not been patented or made the subject of an inventor's certificate issued before the date of this application in any country foreign to the United States of America on any application filed by me or my legal representatives or assigns more than six months prior to this application.

I hereby claim priority benefits under Title 35, United States Code §119(E)(1) of any United States provisional application(s) for patent as indicated below and have also identified below any application for patent on this invention having a filing date before that of the application for patent on which priority is claimed:

<u>Application No.</u>	<u>Date of Filing (day/month/year)</u>	<u>Priority Claimed</u>
60/117,806	01/29/1999	Yes <u>X</u> No <u> </u>

I hereby appoint the following attorneys and agents to prosecute that application and to transact all business in the Patent and Trademark Office connected therewith and to file, to prosecute and to transact all business in connection with all patent applications directed to the invention:

Thomas P. McCracken, Reg. No. 38,548

Address all correspondence to: Thomas P. McCracken at:

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This appointment, including the right to delegate this appointment, shall also apply to the same extent to any proceedings established by the Patent Cooperation Treaty.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under § 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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